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APPLICATION NO.	FII	LING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/051,345	10/051,345 01/18/2002		01/18/2002 Yuen Kai Fung		D6087D 9974	
7.	590	02/08/2005		EXAM	INER	
Dr. Benjamin				ZARA,	JANE J	
Adler & Assoc 8011 Candle La				ART UNIT	PAPER NUMBER	
Houston, TX				1635		
				DATE MAILED: 02/08/200	5	

Please find below and/or attached an Office communication concerning this application or proceeding.

		Angli-Al No	1 A 1! 4/->		
		Application No.	Applicant(s)		
		10/051,345	FUNG ET AL.		
	Office Action Summary	Examiner	Art Unit		
		Jane Zara	1635		
Period fo	The MAILING DATE of this communication app or Reply	ears on the cover sheet with the	correspondence address		
THE - Exte after - If the - If NO - Failt Any	ORTENED STATUTORY PERIOD FOR REPLY MAILING DATE OF THIS COMMUNICATION. nsions of time may be available under the provisions of 37 CFR 1.13 SIX (6) MONTHS from the mailing date of this communication. e period for reply specified above is less than thirty (30) days, a reply period for reply is specified above, the maximum statutory period we are to reply within the set or extended period for reply will, by statute, reply received by the Office later than three months after the mailing ed patent term adjustment. See 37 CFR 1.704(b).	36(a). In no event, however, may a reply be to within the statutory minimum of thirty (30) da will apply and will expire SIX (6) MONTHS fror cause the application to become ABANDON	mely filed ys will be considered timely. n the mailing date of this communication. ED (35 U.S.C. § 133).		
Status					
1)[🛛	Responsive to communication(s) filed on 12 No	ovember 2004.			
		action is non-final.			
3)□	Since this application is in condition for allowar	nce except for formal matters, p	osecution as to the merits is		
	closed in accordance with the practice under E	x parte Quayle, 1935 C.D. 11, 4	53 O.G. 213.		
Disposit	ion of Claims				
5)	 4) Claim(s) 4-7 and 11-13 is/are pending in the application. 4a) Of the above claim(s) is/are withdrawn from consideration. 5) Claim(s) is/are allowed. 6) Claim(s) 4-7 and 11-13 is/are rejected. 7) Claim(s) is/are objected to. 				
8)□	Claim(s) are subject to restriction and/or	r election requirement.			
Applicat	ion Papers				
-	The specification is objected to by the Examine The drawing(s) filed on is/are: a) accomplicant may not request that any objection to the	epted or b) objected to by the drawing(s) be held in abeyance. Se	ee 37 CFR 1.85(a).		
11)	Replacement drawing sheet(s) including the correct The oath or declaration is objected to by the Ex				
Priority (under 35 U.S.C. § 119				
 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of: 1. Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received. 					
Attachmen	ıt(s)	_			
2) Notice 3) Infor	ce of References Cited (PTO-892) ce of Draftsperson's Patent Drawing Review (PTO-948) mation Disclosure Statement(s) (PTO-1449 or PTO/SB/08) or No(s)/Mail Date	4) ☐ Interview Summar Paper No(s)/Mail I 5) ☐ Notice of Informal 6) ☑ Other: <u>NCBI searc</u>	Date Patent Application (PTO-152)		

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DETAILED ACTION

This Office action is in response to the communication filed

Claims 4-7 and 11-13 are pending in the instant application.

The text of those sections of Title 35, U.S. Code not included in this action can be found in a prior Office action.

Response to Arguments and Amendments

Withdrawn Rejections

Any rejections not repeated in this Office action are hereby withdrawn.

Maintained Rejections

The specification is objected to under 35 USC § 112, first paragraph as failing to provide an enabling disclosure for the claimed invention for the same reasons of record set forth in the Office action mailed 5-6-04.

Claims 4-15 are rejected under 35 U.S.C. 112, first paragraph, as based on a disclosure which is not enabling. Features and steps critical or essential to the practice of the invention, but not included in the claim(s) is not enabled by the disclosure for the same reasons of record set forth in the Office action mailed 5-6-04. See *In re Mayhew*, 527 F.2d 1229, 188 USPQ 356 (CCPA 1976).

Applicant's arguments filed 11-12-04 have been fully considered but they are not persuasive. Applicants argue that the claimed invention is adequately described in the disclosure and therefore no deposit is required for the instantly claimed vectors. Contrary to Applicant's assertions, the vectors claimed comprise various and multiple promoters which are to drive the concerted

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expression of the numerous and varied cassettes within the vectors, each of the cassettes in turn comprising many components including competing DNA-binding domains, leucine zipper domains, transactivation domains, IRES sequences, nuclear localization signals, antisense sequences and dominant negative sequences which are directed to corresponding sequences located at other parts of the vector. It would require undue experimentation beyond that taught in the instant disclosure to synthesize the vectors claimed whereby the promoters drive the concerted or orchestrated expression of all downstream components included in the vectors. Inadequate guidance is provided in the instant disclosure regarding the sequences comprising each of the promoters claimed that successfully drive the concerted expression of these various vector components, as well as the portions of the IRES and nuclear localization signals that together provide for functional expression vectors. Therefore vectors pRIBS-X and pRIPS-X are required material for the compositions claimed.

Claims 4-7 and 11-13 are rejected under 35 U.S.C. 112, first paragraph, because the specification, for the same reasons of record set forth in the Office action mailed 5-6-04. The claimed vectors comprise therapeutic genes and no in vivo enablement has been shown for these expression vectors in combination with any therapeutic gene. Therefore the enablement rejection is maintained.

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New Rejections

Claim Rejections - 35 USC § 112

The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

Claims 4-7 and 11-13 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

In claim 4, line 6 (section (a)), it is unclear what amino acid sequence the parenthesized amino acid sequences 1-147 are referring to (e.g. since SEQ ID NO: 1 appears to adequately describe the nucleotide sequences, it seems unnecessary to list amino acid sequence numbers of an undisclosed amino acid SEQ ID NO).

In claim 4, line 7, it is unclear what is meant by the term "fused to" (e.g. indicating operable linkage between the various vector components would be remedial).

In claim 4, line 8 (section (a)), it is unclear what amino acid sequence the parenthesized amino acid sequences 8-112 of Max are referring to (e.g. since SEQ ID NO: 2 appears to adequately describe the nucleotide sequences, it seems unnecessary to list amino acid sequence numbers of an undisclosed amino acid SEQ ID NO).

In claim 4, line 14, it is unclear whether "Gal-DBD" consists of SEQ ID NO:

1, or more amino acids of the DNA-binding domain of yeast GAL4 protein (e.g. see lines 6-7 of claim 4).

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In claim 4, line 8, the term "followed by" is vague (e.g. indicating operable linkage between the various vector components would be remedial).

In claim 4, lines 12-13, it is unclear whether the "sequence complementary to the Gal-DBD-mx sequence" is complementary to all or part of the Gal-DBD-mx sequence.

In claim 4, lines 17-19, it is unclear whether SEQ ID NO: 3 encodes Gal4 alone, or Gal4 in combination with the nuclear localization signal.

Generally in claim 4, it is unclear whether or not all of the component parts of the vector are operably linked.

In claim 4, lines 21-23, it is unclear whether or not SEQ ID NO: 5 encodes the helix loop helix leucine zipper domain of c-Myc and SV40 poly, or only the leucine zipper domain.

In claim 4, line 30, the term "gene X" is vague and unclear (e.g. replacing this with – a gene of interest—would be remedial).

In claim 4, lines 32-33, it is unclear whether the antisense TET-ON sequence is fully or partially complementary to SEQ ID NO: 8.

In claim 7, line 2, the term "gene X" is vague and unclear (e.g. replacing it with the term – gene of interest—would be remedial).

In claim 11, line 6, it is unclear what amino acid sequence the parenthesized amino acid sequences 1-147 are referring to (e.g. since SEQ ID NO: 1 appears to adequately describe the nucleotide sequences, it seems

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unnecessary to list amino acid sequence numbers of an undisclosed amino acid SEQ ID NO).

In claim 11, line 6, there appears to be missing words (e.g. inserting –of the—before "DNA-binding" would be remedial).

Generally in claim 11, it is unclear whether or not all of the component parts of the vector are operably linked.

In claim 11, line 8, it is unclear what amino acid sequence the parenthesized amino acid sequences 8-112 of Max are referring to (e.g. since SEQ ID NO: 2 appears to adequately describe the nucleotide sequences, it seems unnecessary to list amino acid sequence numbers of an undisclosed amino acid SEQ ID NO).

In claim 11, line 8, the term "followed by" is vague (replacing this with – operably linked to—would be remedial).

In claim 11, lines 12-13, it is unclear whether the complementary sequence consists of all or a portion of Gal-DBD-mx.

In claim 11, line 14, it is unclear what sequence comprises the Gal-DBD (e.g. is this SEQ ID NO: 1?).

In claim 11, lines 21-23, it is unclear whether SEQ iD NO: 5 encodes the leucine zipper domain as well as SV40 polyA or only the leucine zipper domain.

In claim 11, line 30, the term "gene X" is vague and unclear (e.g. replacing this with – a gene of interest—would be remedial).

In claim 11, lines 32-34, it is unclear whether the TET-ON antisense is complementary to all or a portion of SEQ ID NO: 8.

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In claim 13, line 2, the term "gene X" is vague and unclear (e.g. replacing it with the term – gene of interest—would be remedial).

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

Claims 4-7 and 11-13 are rejected under 35 U.S.C. 112, first paragraph, as failing to comply with the written description requirement. The claim(s) contains subject matter which was not described in the specification in such a way as to reasonably convey to one skilled in the relevant art that the inventor(s), at the time the application was filed, had possession of the claimed invention.

The claims are drawn to vectors comprising multiple promoters which work in an orchestrated manner to drive expression of a gene of interest. These promoters include any c-erbB2, GAPp-ptet, Egr-1, whey acidic protein, stromelysin 3, pProbasin, or prostate specific antigen promoter. The specification and claims do not adequately describe the broad genera comprising c-erbB2, GAPp-ptet, Egr-1, whey acidic protein, stromelysin 3, pProbasin, or prostate specific antigen promoters. The specification and claims do not indicate the concise features or attributes shared by each of these genera comprising functional c-erbB2, GAPp-ptet, Egr-1, whey acidic protein, stromelysin 3, pProbasin, or prostate specific antigen promoters which drive the expression of the vectors and corresponding components claimed. The scope of the claims

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includes numerous structural variants for each of the genera comprising a particular promoter, and sequence variations exist within each promoter genus, depending on the species from which the promoters are obtained, and/or depending on the molecular source of the promoter obtained from a particular species (see the enclosed NCBI searches of the various promoters). The genera are highly variant because a significant number of structural differences between members of a given genus is permitted. The specification fails to teach or adequately describe a representative number of species in each genus. And because the genera are highly variant, the description provided is insufficient. One of skill in the art would reasonably conclude that the disclosure fails to provide a representative number of species to describe the various genera claimed. Thus, Applicant was not in possession of the claimed genera.

Conclusion

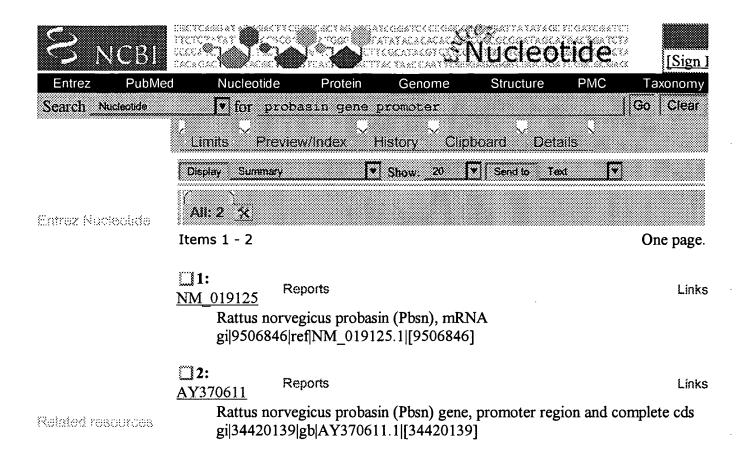
Certain papers related to this application may be submitted to Art Unit 1635 by facsimile transmission. The faxing of such papers must conform with the notices published in the Official Gazette, 1156 OG 61 (November 16, 1993) and 1157 OG 94 (December 28, 1993) (see 37 C.F.R. § 1.6(d)). The official fax telephone number for the Group is **703-872-9306**. NOTE: If Applicant *does* submit a paper by fax, the original signed copy should be retained by applicant or applicant's representative. NO DUPLICATE COPIES SHOULD BE SUBMITTED so as to avoid the processing of duplicate papers in the Office.

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Any inquiry concerning this communication or earlier communications from the examiner should be directed to **Jane Zara** whose telephone number is **(571) 272-0765.** If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, John LeGuyader, can be reached on (571) 272-0760. Any inquiry regarding this application should be directed to the patent analyst, Katrina Turner, whose telephone number is (571) 272-0564. Any inquiry of a general nature or relating to the status of this application should be directed to the Group receptionist whose telephone number is (703) 308-0196.

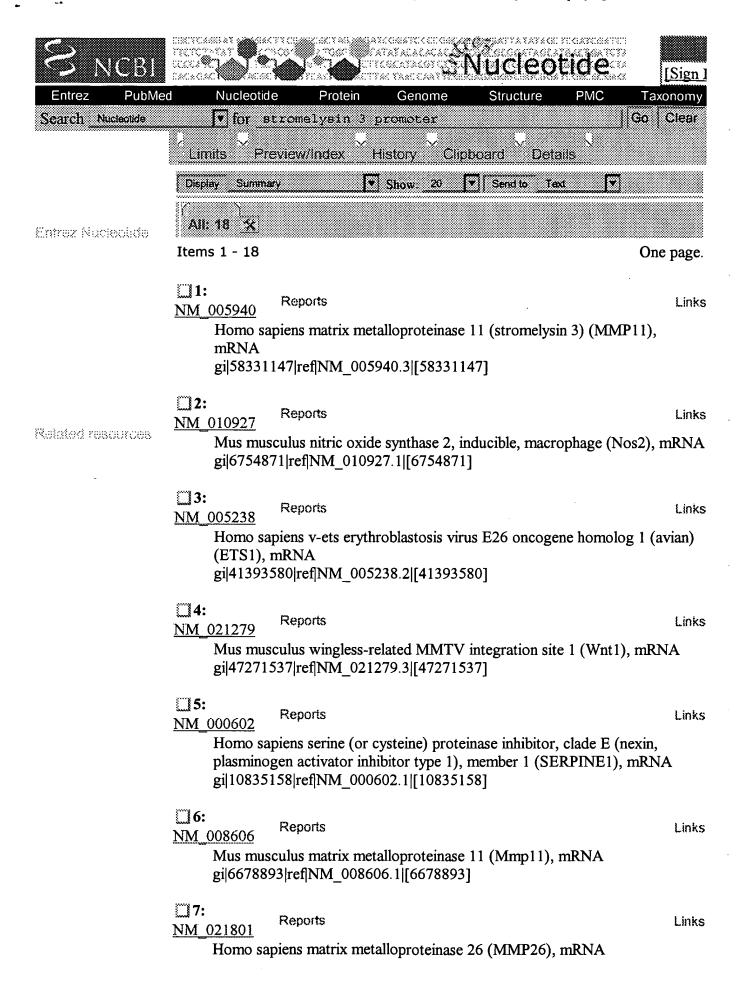
Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

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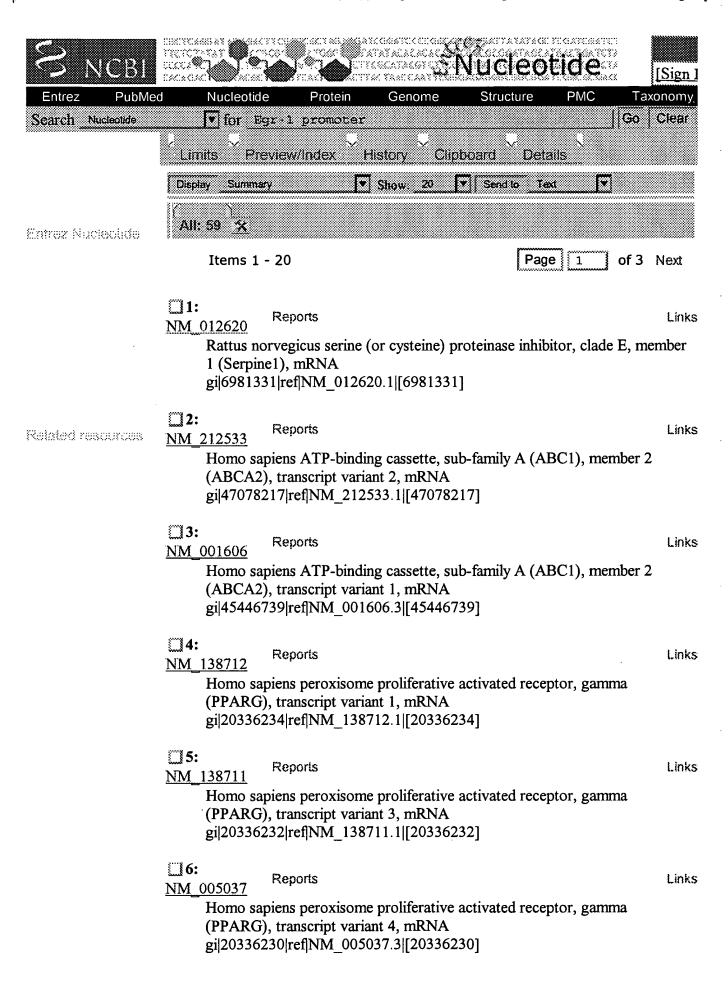
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یه ده: NM	002422	Reports	Links
		piens matrix metalloproteinase 3 (stromelysin 1, progelatinase), mRNA	
	` //	.803 ref NM_002422.2 [13027803]	
□9: NM	002421	Reports	Links
	Homo say	piens matrix metalloproteinase 1 (interstitial collagenase) (MM	P1),
	gi 130277	798 ref NM_002421.2 [13027798]	
[]10 NM): 021964	Reports	Links
		piens zinc finger protein 148 (pHZ-52) (ZNF148), mRNA 035 ref NM_021964.1 [11415035]	
11 NC0	: 005027	Reports	Links
		sp. 1, complete genome 666 ref NC_005027.1 [32470666]	
12 CD20	: 68601	Reports	Links
	TR:P975	x1 Hydra EST -III Hydra magnipapillata cDNA 3' similar to 68 P97568 STROMELYSIN-3 PRECURSOR;, MRNA seque 428 gb CD268601.1 [31056428]	ence
13	: <u>X84664</u>	- "	Links
	-	s stromelysin-3 gene 6 emb X84664.1 HSSTROM3[984746]	
14 AR18	: 32516	Reports	
	-	e 3 from patent US 6338944 723 pat US 6338944 3 gb AR182516.1 [20225723]	
□15 AF29		Reports	Links
		culus stromelysin-3 (Mmp11) gene, promoter and partial cds 608 gb AF297645.1 AF297645[10280608]	
□16 AR04	: 19980	Reports	
	-	3 from patent US 5824794 72 pat US 5824794 3 gb AR049980.1 AR049980[5971972]	

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35-28 pags 18-appg



17: NM 015869	Reports	Links
(PPARG	apiens peroxisome proliferative activated receptor, gamma b), transcript variant 2, mRNA 228 ref NM_015869.2 [20336228]	
	Reports apiens Fas ligand (TNF superfamily, member 6) (FASLG), mRN 28 ref NM_000639.1 [4557328]	Links IA
LIN-21, (mab-5)	Reports abditis elegans male ABnormal MAB-5, abnormal cell LINeage Homeobox C member, required for cell differentiation (22.4 kI complete mRNA 827 ref NM_066294.2 [25144827]	
disease)	Reports Apiens transforming growth factor, beta 1 (Camurati-Engelmann (TGFB1), mRNA 1974 ref NM_000660.2 [56605974]	Links
	Reports apiens vascular endothelial growth factor (VEGF), mRNA 563 ref NM_003376.3 [30172563]	Links
	Reports cetus auratus glucagon gene, promoter region 040 gb AY842856.1 [56609040]	Links
system)	Reports appiens CD44 antigen (homing function and Indian blood group (CD44), transcript variant 5, mRNA [942 ref NM_001001392.1 [48255942]	Links
system)	Reports apiens CD44 antigen (homing function and Indian blood group (CD44), transcript variant 4, mRNA 6940 ref NM_001001391.1 [48255940]	Links
☐15: NM_00100139	00 Reports	Links

gi|48255938|ref|NM 001001390.1|[48255938] **16:** Reports Links NM 001001389 Homo sapiens CD44 antigen (homing function and Indian blood group system) (CD44), transcript variant 2, mRNA gi|48255936|ref|NM 001001389.1|[48255936] **17:** Reports Links NM 000610 Homo sapiens CD44 antigen (homing function and Indian blood group system) (CD44), transcript variant 1, mRNA gi|48255934|ref|NM 000610.3|[48255934] **18:** Reports Links NM 058197 Homo sapiens cyclin-dependent kinase inhibitor 2A (melanoma, p16, inhibits CDK4) (CDKN2A), transcript variant 3, mRNA gi|47132607|ref|NM_058197.2|[47132607] **19:** Reports Links NM 000077 Homo sapiens cyclin-dependent kinase inhibitor 2A (melanoma, p16, inhibits CDK4) (CDKN2A), transcript variant 1, mRNA gi|47132606|ref|NM_000077.3|[47132606]

Homo sapiens CD44 antigen (homing function and Indian blood group

system) (CD44), transcript variant 3, mRNA

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NM 058195

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Homo sapiens cyclin-dependent kinase inhibitor 2A (melanoma, p16, inhibits CDK4) (CDKN2A), transcript variant 4, mRNA gi|47132605|ref|NM 058195.2|[47132605]

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Summary

Reports

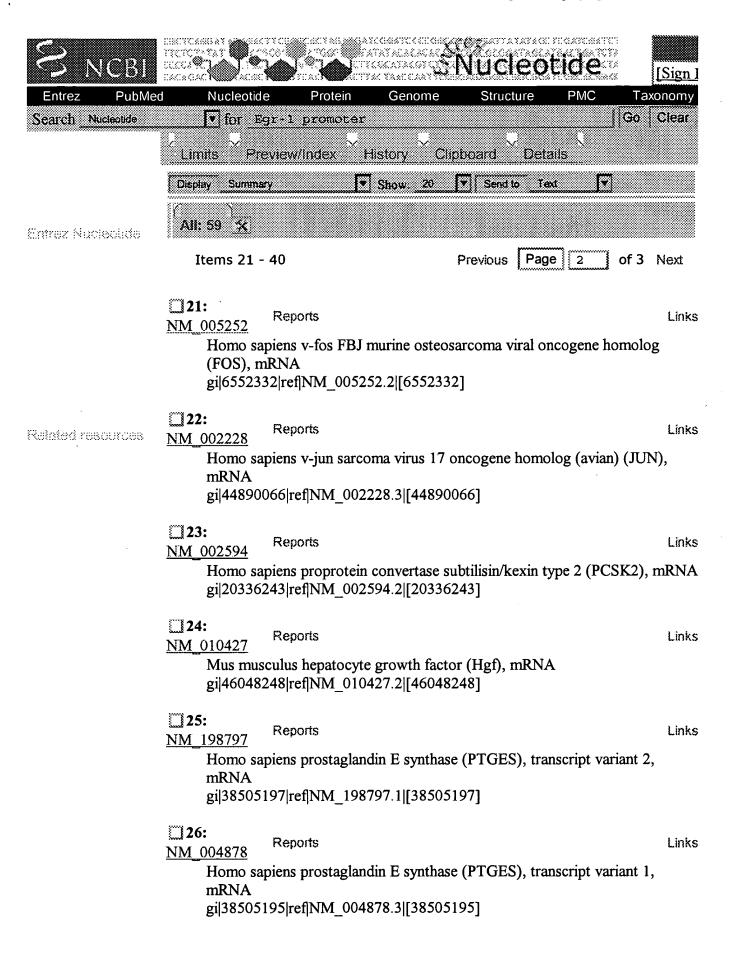
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27: Reports	Links
NM_000898 Homo sapiens monoamine oxidase B (MAOB), nuclear gene encoding mitochondrial protein, mRNA gi 38202206 ref NM_000898.3 [38202206]	
NM_010177 Reports Mus musculus tumor necrosis factor (ligand) superfamily, member 6 (Tnfsf6), mRNA gi 31981778 ref NM_010177.2 [31981778]	Links
29: NM_007913 Reports Mus musculus early growth response 1 (Egr1), mRNA gi 24475900 ref NM_007913.2 [24475900]	Links
NM_000602 Homo sapiens serine (or cysteine) proteinase inhibitor, clade E (nexin, plasminogen activator inhibitor type 1), member 1 (SERPINE1), mRNA gi 10835158 ref NM_000602.1 [10835158]	Links
MM_004864 Homo sapiens growth differentiation factor 15 (GDF15), mRNA gi 4758935 ref NM_004864.1 [4758935]	Links
T32: NM_002930 Homo sapiens Ras-like without CAAX 2 (RIT2), mRNA gi 4506532 ref NM_002930.1 [4506532]	Links
Mus musculus gonadotropin releasing hormone 2 (Gnrh2), mRNA gi 51093848 ref NM_008145.1 [51093848]	Links
Tak: NM_001964 Homo sapiens early growth response 1 (EGR1), mRNA gi 31317226 ref NM_001964.2 [31317226]	Links
MM_013261 Reports Homo sapiens peroxisome proliferative activated receptor, gamma, coactivator 1, alpha (PPARGC1A), mRNA pil29570796lreflNM_013261.2l[29570796]	Links

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Items 21 -	40	Previous Pa	age 2 of 3	Next
variant 1,	piens mitogen-activated prote mRNA 528 ref NM_002745.2 [20986	`	IAPK1), transcrip	t
40: NM 002745	Reports			Links
variant 2,	Reports piens mitogen-activated proto mRNA 530 ref NM_138957.1 [20986	`	1APK1), transcrip	Links t
	Reports culus early growth response 1051 ref NM_010118.1 [23956	` • /·	JA	Links
	Reports culus mitogen activated proto 663 ref NM_011949.2 [27370	•	Iapk1), mRNA	Links
Homo sap	piens interleukin 1, beta (IL1 305 ref NM_000576.2 [27894	• •		
□ 36: NM 000576	Reports			Links

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	Items 41 - 59 Previous Page 3 of	f 3
	M 053056 Homo sapiens cyclin D1 (PRAD1: parathyroid adenomatosis 1) (CCND1 mRNA gi 16950654 ref NM_053056.1 [16950654]	Links l),
Related resources	M 033150 Homo sapiens collagen, type II, alpha 1 (primary osteoarthritis, spondyloepiphyseal dysplasia, congenital) (COL2A1), transcript variant 2 mRNA gi 15149478 ref NM_033150.1 [15149478]	Links 2,
	Id3: NM_001844 Homo sapiens collagen, type II, alpha 1 (primary osteoarthritis, spondyloepiphyseal dysplasia, congenital) (COL2A1), transcript variant 1 mRNA gi 15149477 ref NM_001844.3 [15149477]	Links I,
	Reports Rattus norvegicus apurinic/apyrimidinic endonuclease 1 (Apex1), mRNA gi 13162336 ref NM_024148.1 [13162336]	Links
	MM_001275 Reports Homo sapiens chromogranin A (parathyroid secretory protein 1) (CHGA mRNA gi 10800418 ref NM_001275.2 [10800418]	Links
	Mus musculus tumor necrosis factor (Tnf), mRNA gi 7305584 ref NM_013693.1 [7305584]	Links

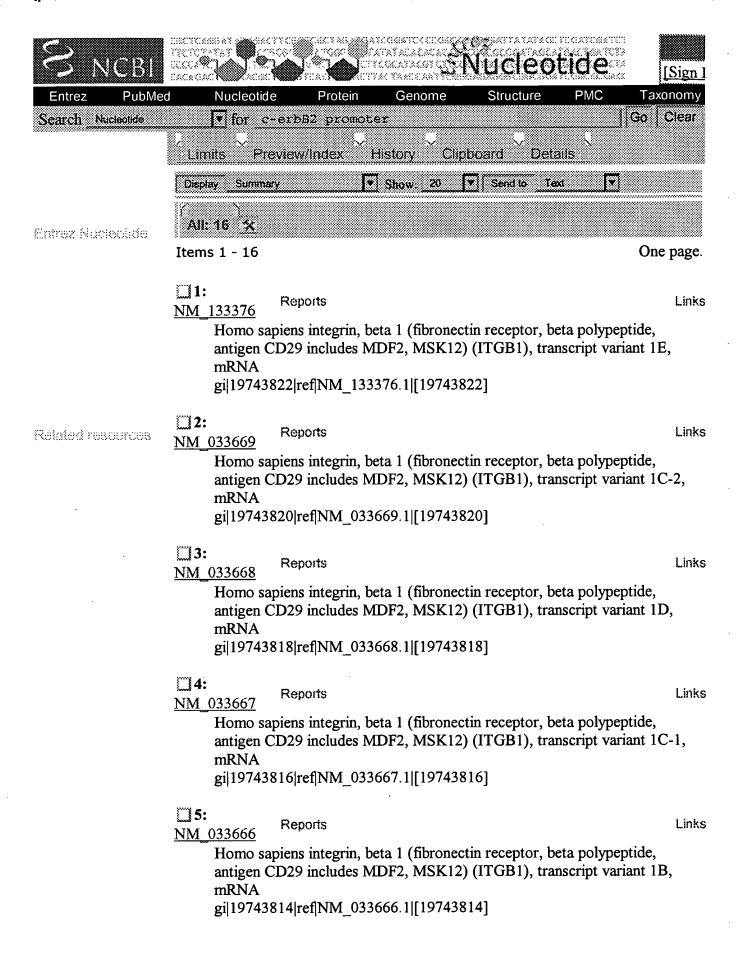
147: NM 011146	Reports	Links
	culus peroxisome proliferator activated receptor gamma (Ppara	g),
	37 ref NM_011146.1 [6755137]	
AY 630403	Reports	Links
	culus IL-2 receptor beta-chain gene, promoter region and 5'U7 206 gb AY630403.1 [49338206]	'R
149: AX814795	Reports	Links
-	e 1 from Patent WO03064465 989 emb AX814795.1 pat WO 03064465 1[39103989]	
50: NC_003283	Reports	Links
	abditis elegans chromosome V, complete sequence 584 ref NC_003283.3 [32967584]	
51: AY142704	Reports	Links
	allus transcription factor CEF-5 gene, complete cds 434 gb AY142704.1 [23380434]	
52: AB083340	Reports	Links
	culus mPGES gene, promoter, partial sequence 892 dbj AB083340.1 [21327892]	
53: AY029236	Reports	Links
Homo sa partial cd		r and
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54: AX009737	Reports	Links
-	e 7 from Patent WO9960142 34 emb AX009737.1 pat WO 9960142 7[9996934]	
55: AX009736	Reports	Links
-	e 6 from Patent WO9960142 33 emb AX009736.1 pat WO 9960142 6[9996933]	
56. X71791	Reports	Links

Rattus norvegicus partial Gdn/Pn-1 gene for glia-derived nexin/protease nexin I, enhancer region gi|9968728|emb|X71791.2|RNGDNPN1[9968728]

AJ245926	Reports		Links
Homo region	sapiens Egr-1 gene	for early growth response factor	r-1, promoter
gi 6688	3175 emb AJ245926	5.1 HSA245926[6688175]	
58: <u>U735</u>		outons 2 (DC2) come over 1 and	Links
	-	ertase 2 (PC2) gene, exon 1 and HSU73595[2623384]	partial cus
59: X126	17 Reports		Links
region	mgEgr-1 gene for a	mitogen inducible zinc finger pro	otein 5'-flanking
5 13000) O O O O O O O O O		
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16: NM 002211 Reports	Links
Homo sapiens integrin, beta 1 (fibronectin receptor, beta polypeptide, antigen CD29 includes MDF2, MSK12) (ITGB1), transcript variant 1A, mRNA	
gi 19743812 ref NM_002211.2 [19743812]	
<u>NM_001005862</u>	Links
Homo sapiens v-erb-b2 erythroblastic leukemia viral oncogene homolog neuro/glioblastoma derived oncogene homolog (avian) (ERBB2), transcr variant 2, mRNA gi 54792097 ref NM_001005862.1 [54792097]	•
18: NM 004448 Reports	Links
Homo sapiens v-erb-b2 erythroblastic leukemia viral oncogene homolog neuro/glioblastoma derived oncogene homolog (avian) (ERBB2), transcr variant 1, mRNA gi 54792095 ref NM 004448.2 [54792095]	
19:	
NM_000546 Reports	Links
Homo sapiens tumor protein p53 (Li-Fraumeni syndrome) (TP53), mRN gi 8400737 ref NM_000546.2 [8400737]	Α
10: Reports NM 002203	Links
Homo sapiens integrin, alpha 2 (CD49B, alpha 2 subunit of VLA-2 receptor), mRNA gi 6006008 ref NM_002203.2 [6006008]	otor)
T11: Reports	Links
NM_002467	Lanco
Homo sapiens v-myc myelocytomatosis viral oncogene homolog (avian) (MYC), mRNA gi 31543215 ref NM_002467.2 [31543215]	
12: NM 002012 Reports	Links
Homo sapiens fragile histidine triad gene (FHIT), mRNA gi 4503718 ref NM_002012.1 [4503718]	
13: NM 010152 Reports	Links
Mus musculus v-erb-b2 erythroblastic leukemia viral oncogene homolog neuro/glioblastoma derived oncogene homolog (avian) (Erbb2), transcrip variant 2, mRNA gi 54873611 ref NM_010152.2 [54873611]	

14: Reports Links NM 001003817 Mus musculus v-erb-b2 erythroblastic leukemia viral oncogene homolog 2, neuro/glioblastoma derived oncogene homolog (avian) (Erbb2), transcript variant 1, mRNA gi|54873609|ref|NM_001003817.1|[54873609] **15**: Reports Links NM_053056 Homo sapiens cyclin D1 (PRAD1: parathyroid adenomatosis 1) (CCND1), **mRNA** gi|16950654|ref|NM_053056.1|[16950654] **16:** <u>X56495</u> Reports Links H.sapiens DNA for the upstream regulatory region of the c-erbB2 gene gi|29880|emb|X56495.1|HSCERBB2[29880]

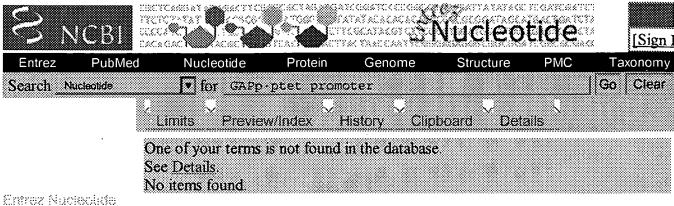
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Summary

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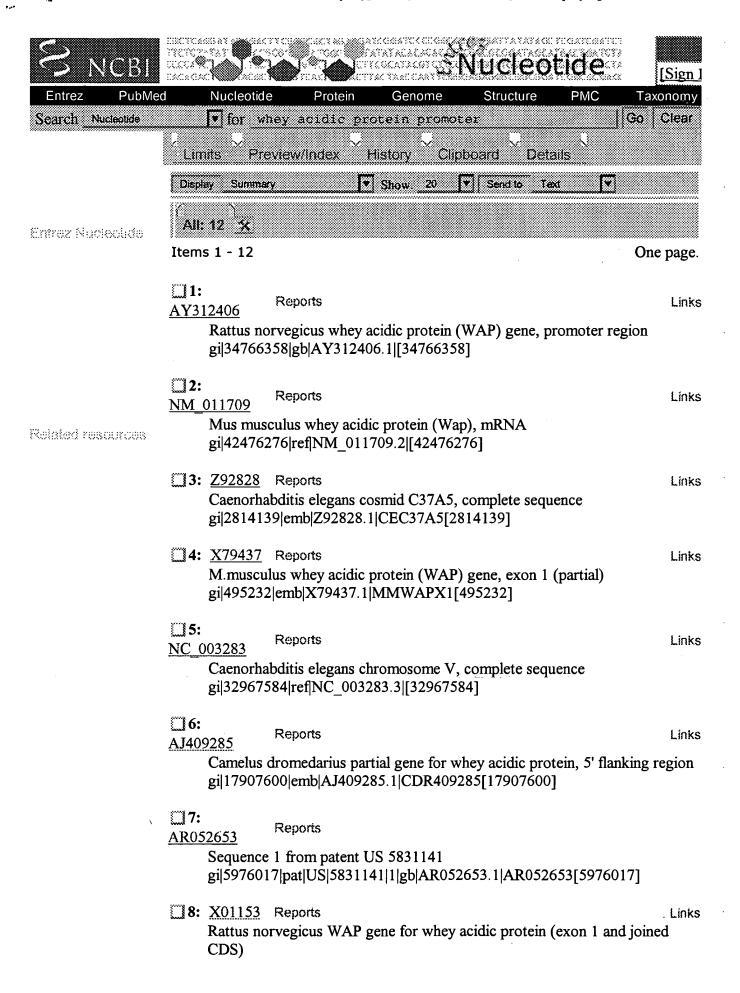
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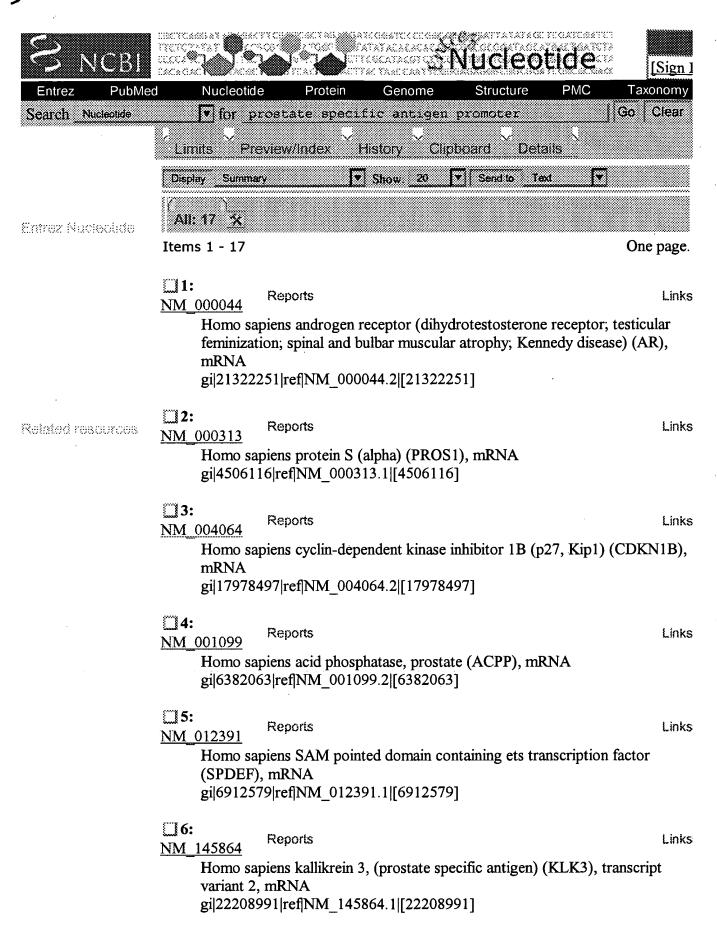
gi|57492|emb|X01153.1|RNWAP1[57492]

9: AA40) <u>8564</u>	Reports					Links
	musculus		C0030B08	3', MRNA se	al cone cDNA equence	A library Mu	S
10	J0068D10 promoter	Mouse 3.5-dp	Mus muscu sequence	lus whey aci	musculus cD dic protein (\		Links
11:	: <u>U38816</u>	Reports					Links
		_	_	(Wap) gene, 316[1215721	, promoter an	nd complete	cds
12 :	: <u>L.21193</u>	Reports					Links
		ney acidic pro gb L21193.		romoter regi P[309534]	ion		
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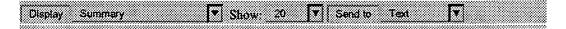
□7: NM	001648	Reports	Links
	variant 1,	piens kallikrein 3, (prostate specific antigen) (KLK3), transcription mRNA 090 ref NM_001648.2 [22208990]	t
☐8: NM	-	Reports piens macrophage migration inhibitory factor ation-inhibiting factor) (MIF), mRNA	Links
		84 ref NM_002415.1 [4505184]	
<u></u> 9:	011109	Reports	Links
	_	piens chromosome 19 genomic contig 594 ref NT_011109.15 Hs19_11266[29800594]	
□10 <u>AY28</u>): 33617	Reports	Links
	-	piens KLK3 gene promoter region, partial sequence 297 gb AY283617.1 [32879297]	
11 AY28	: 83616	Reports	Links
	-	piens KLK3 gene promoter region, partial sequence 296 gb AY283616.1 [32879296]	
12 AY28	83615	Reports	Links
		piens KLK3 gene promoter region, partial sequence 295 gb AY283615.1 [32879295]	
13 AY28	8 : 83614	Reports	Links
		piens KLK3 gene promoter region, partial sequence 294 gb AY283614.1 [32879294]	
□14 AY28	l : 83613	Reports	Links
	-	piens KLK3 gene promoter region, partial sequence 293 gb AY283613.1 [32879293]	
15 AY28	5 : 83612	Reports	Links
	-	piens KLK3 gene promoter region, partial sequence 292 gb AY283612.1 [32879292]	
[]16	5: <u>S81389</u> prostate-s	Reports specific antigen/PCPSA {promoter} [human, prostate cancer p	Links atient

isolate, Genomic, 620 nt] gi|1336767|bbm|384056|bbs|176641|gb|S81389.1|S81389[1336767]

17: <u>X92553</u> Reports

Links

H.sapiens aps gene promoter region gi|1262808|emb|X92553.1|HSAPSPROM[1262808]



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